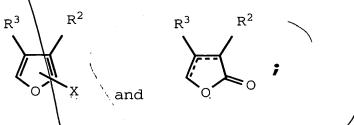
## -- 35. A compound of a formula selected from



wherein X is H or hydroxyl;

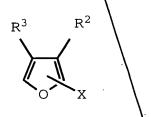
wherein R<sup>2</sup> is a substituent selected from pyridyl, and aryl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower\_alkylsulfonyl, nitro, amino, lower alkylamino, sulfamyl and lower alkylsulfonylamino; and

wherein R<sup>3</sup> is a substituent selected from pyridyl, and aryl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino, lower alkylamino amide, lower alkylsulfonylamino and sulfamyl

provided that at least one of said  $R^2$  and  $R^3$  substituents is substituted with lower alkylsulfonyl or sulfamyl;

or a pharmaceutically-acceptable salt thereof. --

## -- 36. A compound of the \formula



wherein X is H or hydroxyl;

wherein  $R^2$  is a substituent selected from pyridyl, and aryl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy,

lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino lower alkylamino, sulfamyl and lower alkylsulfonylamino; and

wherein R<sup>3</sup> is a substituent selected from pyridyl, and aryl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino, lower alkylamino, amide, lower alkylsulfonylamino and sulfamyl;

provided that at least one of said  $R^2$  and  $R^3$  substituents is substituted with lower alkylsulfonyl or sulfamyl;

or a pharmaceutically-acceptable salt thereof. --

-- 37. Compound of Claim 36 wherein X is H or hydroxyl; wherein R<sup>2</sup> is a substituent selected from pyridyl, naphthyl and phenyl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkyl, lower alkylsulfinyl, lower alkylsulfinyl, nitro, amino, lower alkylamino, sulfamyl and lower alkylsulfonylamino; and

wherein R<sup>3</sup> is a substituent selected from pyridyl, naphthyl and phenyl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino, lower alkylamino, amide, lower alkylsulfonylamino and sulfamyl; or a pharmaceutically-acceptable salt thereof. --

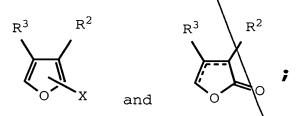
hydroxyl; wherein R<sup>2</sup> is phenyl optionally substituted at a substitutable position with a radical selected from fluoro, chloro, bromo, iodo, methyl, ethyl, isopropyl, tert-butyl, methoxy, ethoxy, butoxy, methylthio, methylsulfinyl, methylsulfonyl, nitro, amino, methylamino, and sulfamyl; and wherein R<sup>3</sup> is phenyl optionally substituted at a substitutable position with a radical selected from fluoro, chloro, bromo, iodo, methyl, ethyl, isopropyl, tert-butyl, methoxy, ethoxy,

butoxy, methylthio, methylsulfinyl, methylsulfonyl, nitro, amino, methylamino, and sulfamyl; or a pharmaceutically-acceptable salt thereof. --

hydroxyl; wherein R<sup>2</sup> is phenyl optionally substituted at a substitutable position with a radical selected from fluoro, chloro, bromo, methyl, ethyl, isopropyl, tert-butyl, methoxy, ethoxy, butoxy, methylsulfonyl, and sulfamyl; and wherein R<sup>3</sup> is phenyl optionally substituted at a substitutable position with a radical selected from fluoro, chloro, bromo, methyl, ethyl, isopropyl, tert-butyl, methoxy, ethoxy, butoxy, methylsulfonyl, and sulfamyl; or a pharmaceutically-acceptable salt thereof. --

-- 40. Compound of claim 39 which is 3-(4-fluorophenyl)-4-(methylsulfonylphenyl) furan, or a pharmaceutically-acceptable salt thereof. --

-- 41. A pharmaceutical composition comprising a therapeutically-effective amount of an antiinflammatory compound, said compound selected from a compound of the formulas



wherein X is H or hydroxyl; wherein each of  $R^2$  and  $R^3$  is a substituent independently selected from aryl and heteroaryl, wherein each of said  $R^2$  and  $R^3$  substituents is optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino, lower alkylamino, sulfamyl and lower

alkylsulfonylamino; provided that at least one of said R<sup>2</sup> and R<sup>3</sup> substituents is substituted with lower alkylsulfonyl or sulfamyl; or a pharmaceutically-acceptable salt thereof. --

-- 42. A pharmaceutical composition comprising a therapeutically-effective amount of an antiinflammatory compound, said compound selected from a compound of the formula

hy h

wherein X is H or hydroxyl;

wherein R<sup>2</sup> is a substituent selected from pyridyl, and aryl optionally substituted at a substitutable position with a radical selected from halo lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino, lower alkylamino sulfamyl and lower alkylsulfonylamino; and

wherein R<sup>3</sup> is a substituent selected from pyridyl, and aryl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino, lower alkylamino, amide, lower alkylsulfonylamino and sulfamyl;

provided that at least one of said  $R^2$  and  $R^3$  substituents is substituted with lower alkylsulfonyl or sulfamyl;

or a pharmaceutically-acceptable salt thereof. --

fluoro, chloro, bromo, iodo, or hydroxyl; wherein X is H, substituent selected from pyridyl, naphthyl and phenyl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino, lower alkylamino, sulfamyl and lower alkylsulfonylamino; and

wherein R<sup>3</sup> is a substituent selected from pyridyl, naphthyl and phenyl optionally substituted at a substitutable position with a radical selected from halo, lower alkyl, lower alkoxy, lower alkylthio, lower alkylsulfinyl, lower alkylsulfonyl, nitro, amino, lower alkylamino, amide, lower alkylsulfonylamino and sulfamyl; or a pharmaceutically-acceptable salt thereof. --

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- -- 44. The composition of Claim 43 wherein X is H or hydroxyl; wherein R<sup>2</sup> is phenyl optionally substituted at a substitutable position with a radical selected from fluoro, chloro, bromo, iodo, methyl, ethyl, isopropyl, tert-butyl, methoxy, ethoxy, butoxy, methylthio, methylsulfinyl, methylsulfonyl, nitro, amino, methylamino, and sulfamyl; and wherein R<sup>3</sup> is phenyl optionally substituted at a substitutable position with a radical selected from fluoro, chloro, bromo, iodo, methyl, ethyl, isopropyl, tert-butyl, methoxy, ethoxy, butoxy, methylthio, methylsulfinyl, methylsulfonyl, nitro, amino, methylamino, and sulfamyl; or a pharmaceutically-acceptable salt thereof. --
- -- 45. The composition of Claim 44 wherein X is H or hydroxyl; wherein R<sup>2</sup> is phenyl optionally substituted at a substitutable position with a radical selected from fluoro, chloro, bromo, methyl, ethyl, isopropyl tert-butyl, methoxy, ethoxy, butoxy, methylsulfonyl, and sulfamyl; and wherein R<sup>3</sup> is phenyl optionally substituted at a substitutable position with a radical selected from fluoro, chloro, bromo, methyl, ethyl, isopropyl, tert-butyl, methoxy, ethoxy, butoxy, methylsulfonyl, and sulfamyl; or a pharmaceutically-acceptable salt thereof. --
- -- 46. The pharmaceutical composition of Claim 42 wherein said compound is 3-(4-fluorophenyl)-4- (methylsulfonylphenyl)furan, or a pharmaceutically acceptable salt thereof. --